STM - Structure Search
1/11/08

10/583,468

=> d ibib abs hitstr 1-3

L4 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2006:15873 CAPLUS
DOCUMENT NUMBER: 144:108216
TITLE: Preparation of amido compounds as inhibitors of 11-β-hydroxysteroid dehydrogenase type 1 (11βHSD1) and antagonists of the mineralocorticoid receptor (MR)
INVENTOR(S): Yao, Wenqing; Xu, Meizhong; Zhang, Colin; Agrios, Konstantinos; Metcalf, Brian; Zhuo, Jincong
PATENT ASSIGNEE(S): Incyte Corporation, USA
SOURCE: PCT Int. Appl., 108 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
WO 2006002349	A1	20060105	WO 2005-US22411		20050623
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW					
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM					
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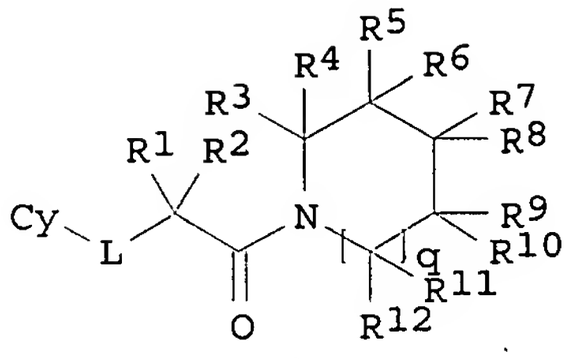
CA 2571258	A1	20060105	CA 2005-2571258		20050623
US 2006009471	A1	20060112	US 2005-159724		20050623
EP 1758582	A1	20070307	EP 2005-762543		20050623
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, LV, MK, YU					
CN 1988908	A	20070627	CN 2005-80020965		20050623
IN 2006KN03601	A	20070615	IN 2006-KN3601		20061201
KR 2007024639	A	20070302	KR 2006-727142		20061222
NO 2007000372	A	20070308	NO 2007-372		20070119
PRIORITY APPLN. INFO.:			US 2004-582556P	P	20040624
			US 2004-639179P	P	20041222
			WO 2005-US22411	W	20050623

OTHER SOURCE(S): MARPAT 144:108216
GI



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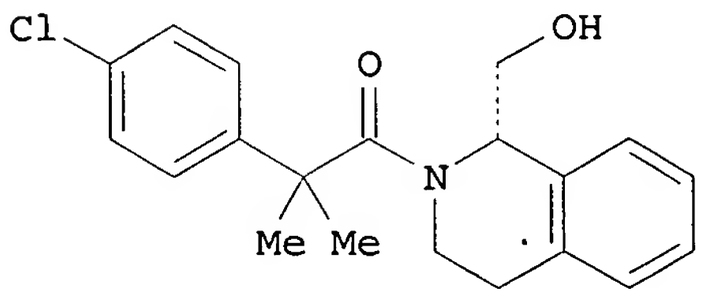


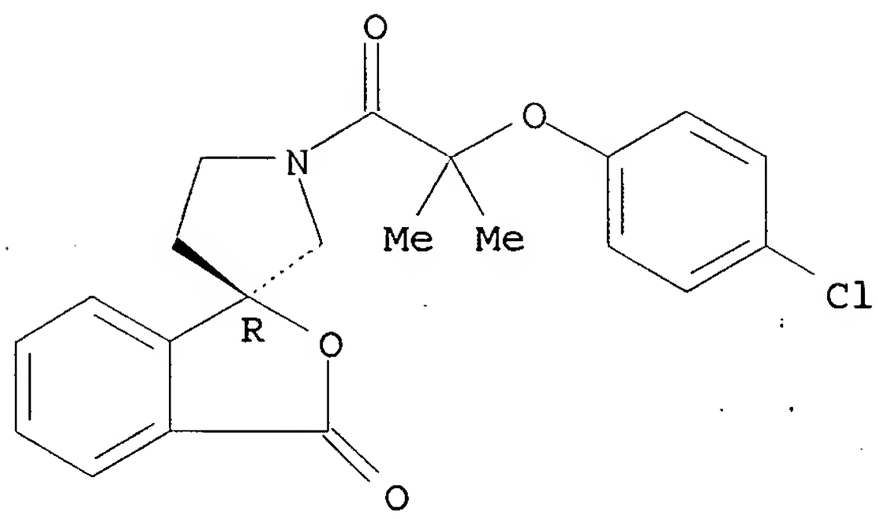
II

AB The title compds. I [Cy = (un)substituted (hetero)aryl, (hetero)cycloalkyl; L = absent, (CR13R14)m, (CR13R14)nO(CR13R14)p, etc.; R1, R2 = (un)substituted alkyl; R3-R12 = H, W1X1Y1Z1; or R3 and R4 together or R5 and R6 together or R7 and R8 together or R9 and R10 together or R11 and R12 together form 4-20 membered cycloalkyl or (un)substituted heterocycloalkyl; or R3 and R12 together or R3 and R10 together or R3 and R8 together or R5 and R10 together or R5 and R10 together or R7 and R12 together form (un)substituted alkylene bridge; R13, R14 = H, halo, alkyl, etc.; W1 = absent, alkylenyl, O, etc.; X1 = absent, alkylenyl, aryl, etc.; Y1 = absent, O, S, etc.; Z1 = H, halo, CN, etc.; m = 1-4; n = 0-3; p = 0-3; q = 0-2; with the provisos] which are inhibitors of 11-β hydroxysteroid dehydrogenase type 1 and antagonists of the mineralocorticoid receptor (MR), were prepared Thus, reacting 2-(4-chlorophenyl)-2-methylpropanoic acid with (1S)-1,2,3,4-tetrahydroisoquinolin-1-ylmethanol in the presence of BOP and N-methylmorpholine in DMF afforded (1S)-II. The compds. I can be useful in the treatment of various diseases associated with expression or activity of 11-β hydroxysteroid dehydrogenase type 1 and/or diseases associated with aldosterone excess. The pharmaceutical composition comprising the compound
I is disclosed.

IT 872985-55-2P 872986-34-0P 872986-36-2P
RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(preparation of amido compds. as inhibitors of 11-β-hydroxysteroid dehydrogenase type 1 (11βHSD1) and antagonists of the mineralocorticoid receptor (MR))

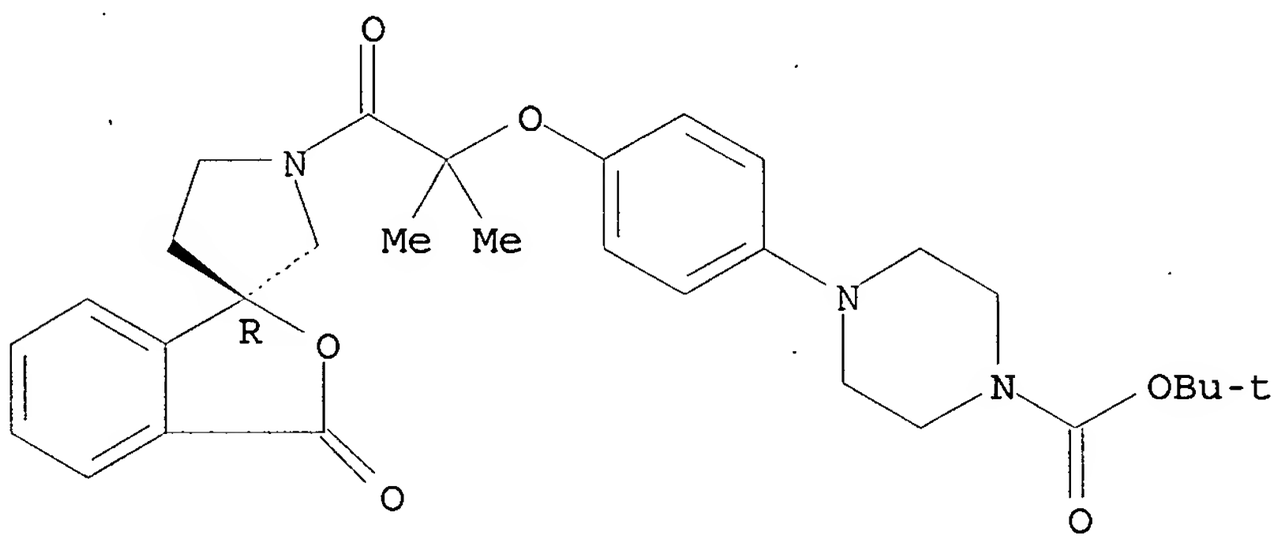
RN 872985-55-2 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(4-chlorophenoxy)-2-methyl-1-oxopropyl]-, (1'R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



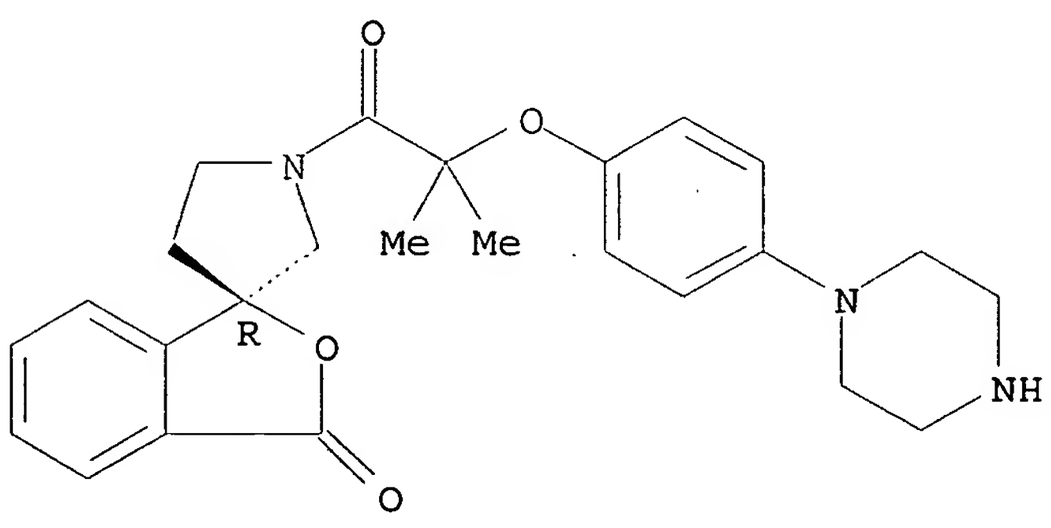
RN 872986-34-0 CAPLUS
CN 1-Piperazinecarboxylic acid, 4-[4-[1,1-dimethyl-2-oxo-2-[(1R)-3-oxospiro[isobenzofuran-1(3H),3'-pyrrolidin]-1'-yl]ethoxy]phenyl]-, 1,1-dimethylethyl ester (CA INDEX NAME)

Absolute stereochemistry.



RN 872986-36-2 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-methyl-1-oxo-2-[4-(1-piperazinyl)phenoxy]propyl]-, (1R)- (CA INDEX NAME)

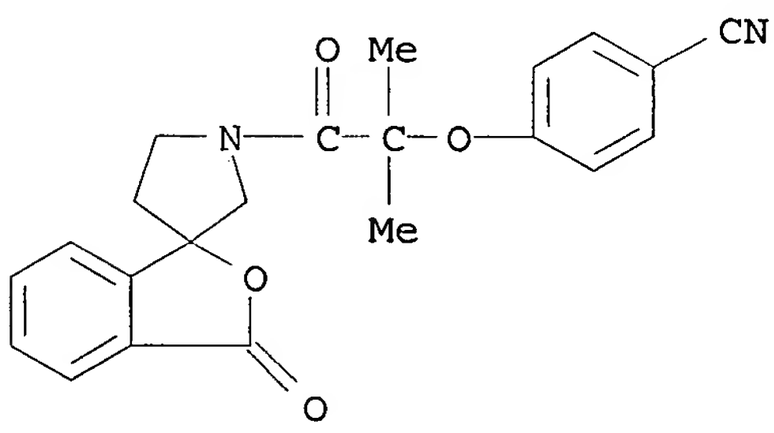
Absolute stereochemistry.



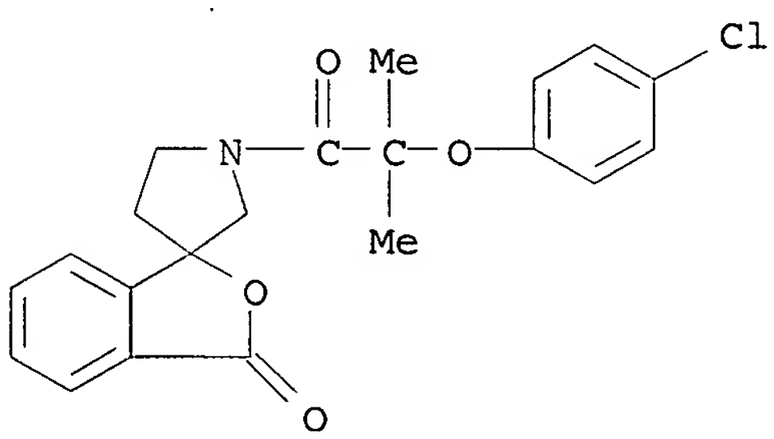
IT 872985-48-3P 872985-49-4P 872985-50-7P
872985-51-8P 872985-52-9P 872985-53-0P
872985-54-1P 872985-56-3P 872985-57-4P
872986-15-7P 872986-19-1P 872986-21-5P
872986-23-7P 872986-25-9P 872986-27-1P
872986-38-4P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU

(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of amido compds. as inhibitors of 11-β-hydroxysteroid dehydrogenase type 1 (11βHSD1) and antagonists of the mineralocorticoid receptor (MR))

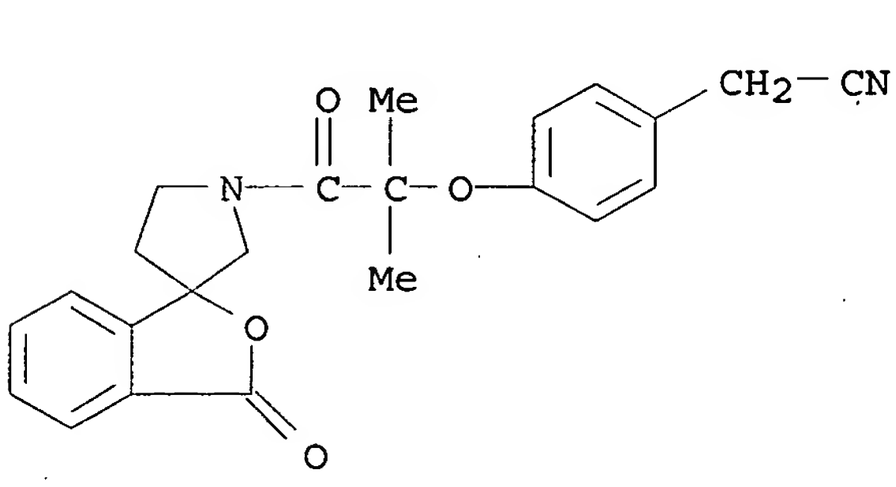
RN 872985-48-3 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(4-cyanophenoxy)-2-methyl-1-oxopropyl]- (9CI) (CA INDEX NAME)



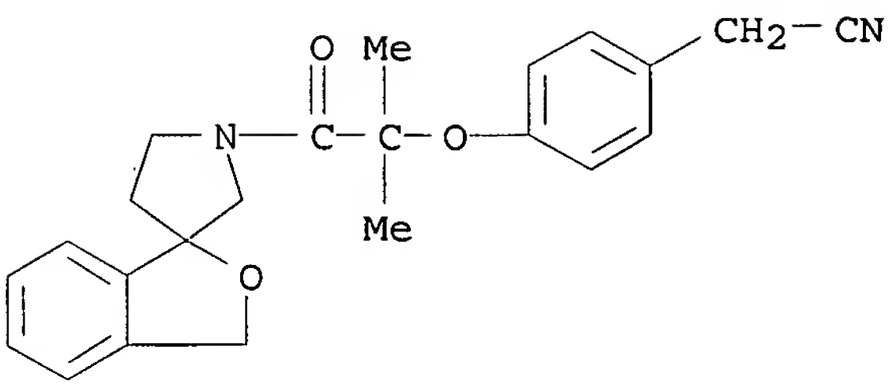
RN 872985-49-4 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(4-chlorophenoxy)-2-methyl-1-oxopropyl]- (CA INDEX NAME)



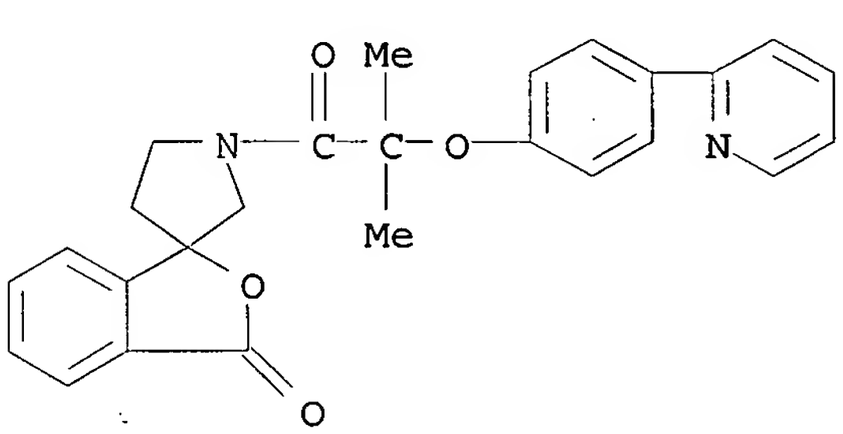
RN 872985-50-7 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-[4-(cyanomethyl)phenoxy]-2-methyl-1-oxopropyl]- (9CI) (CA INDEX NAME)



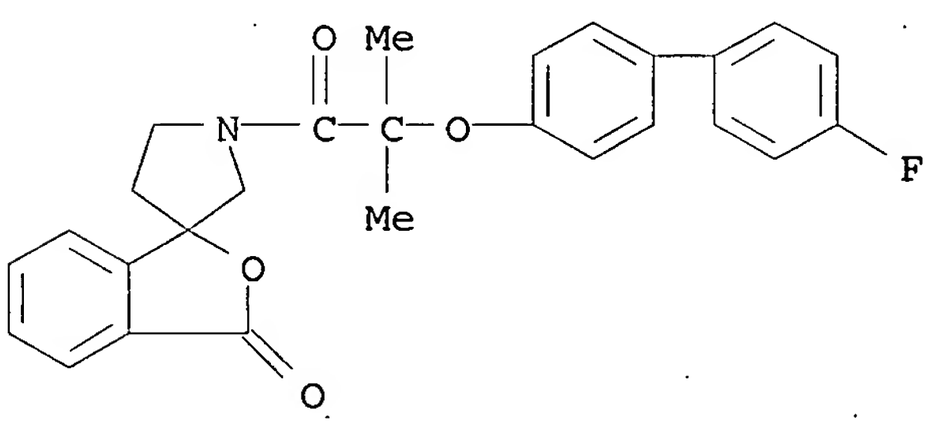
RN 872985-51-8 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidine], 1'-[2-[4-(cyanomethyl)phenoxy]-2-methyl-1-oxopropyl]- (9CI) (CA INDEX NAME)



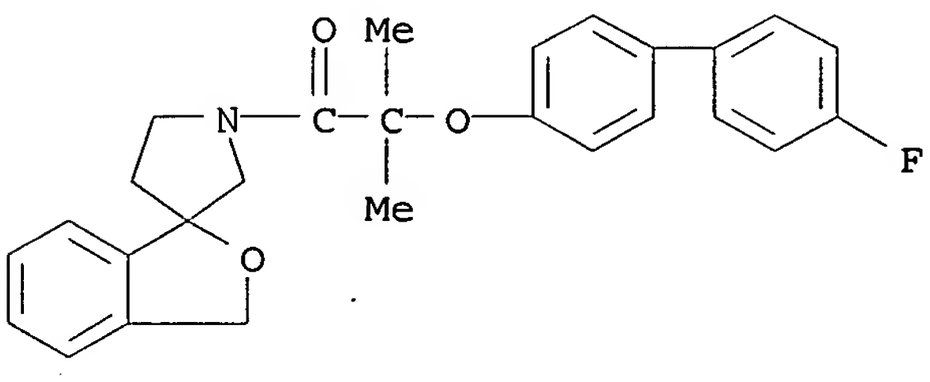
RN 872985-52-9 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-methyl-1-oxo-2-[4-(2-pyridinyl)phenoxy]propyl]- (CA INDEX NAME)



RN 872985-53-0 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-[(4'-fluoro[1,1'-biphenyl]-4-yl)oxy]-2-methyl-1-oxopropyl]- (CA INDEX NAME)

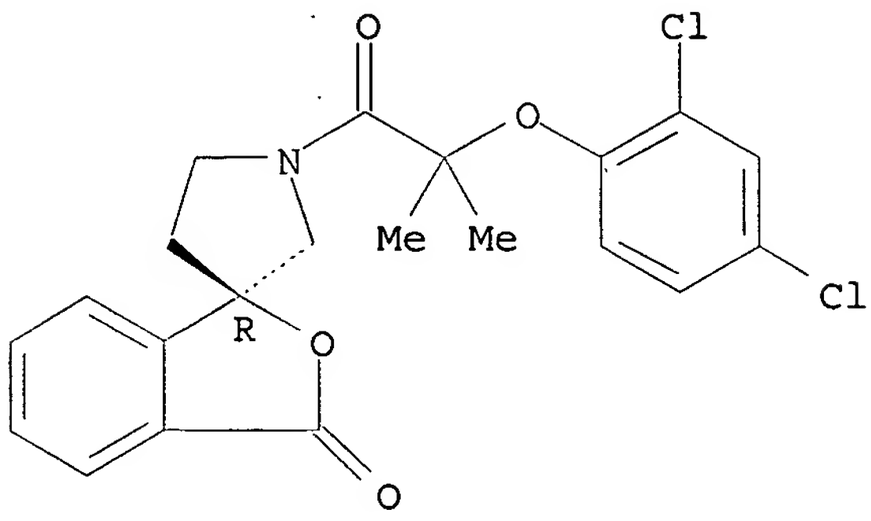


RN 872985-54-1 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidine], 1'-[2-[(4'-fluoro[1,1'-biphenyl]-4-yl)oxy]-2-methyl-1-oxopropyl]- (9CI) (CA INDEX NAME)



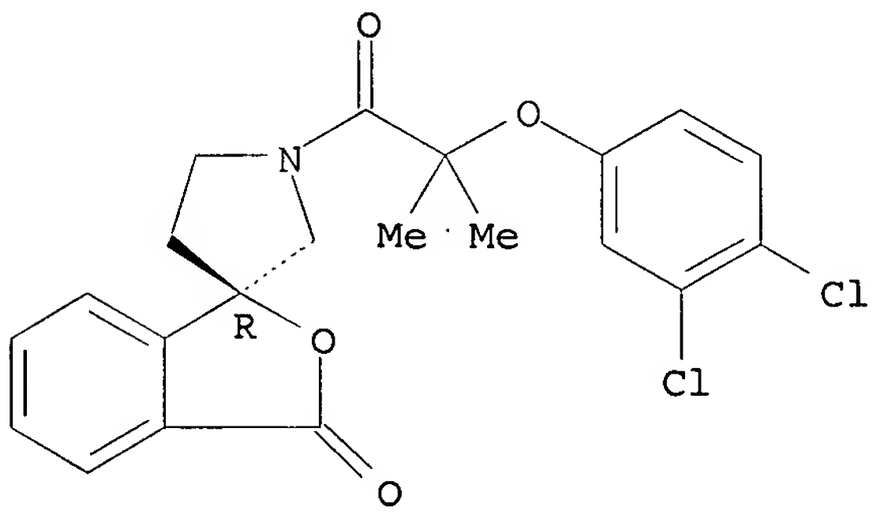
RN 872985-56-3 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(2,4-dichlorophenoxy)-2-methyl-1-oxopropyl]-, (1'R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



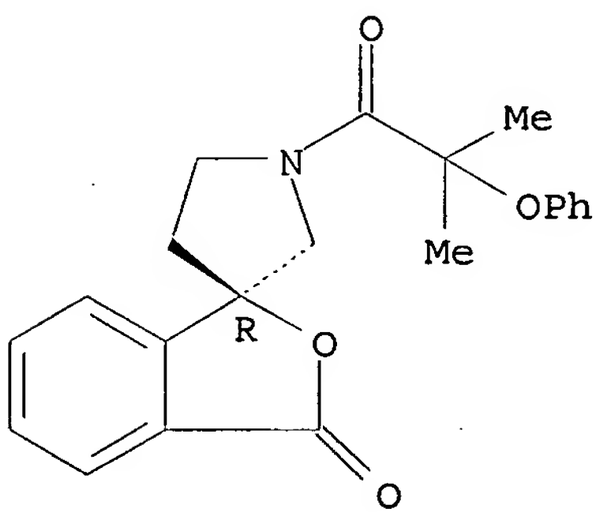
RN 872985-57-4 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(3,4-dichlorophenoxy)-2-methyl-1-oxopropyl]-, (1'R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



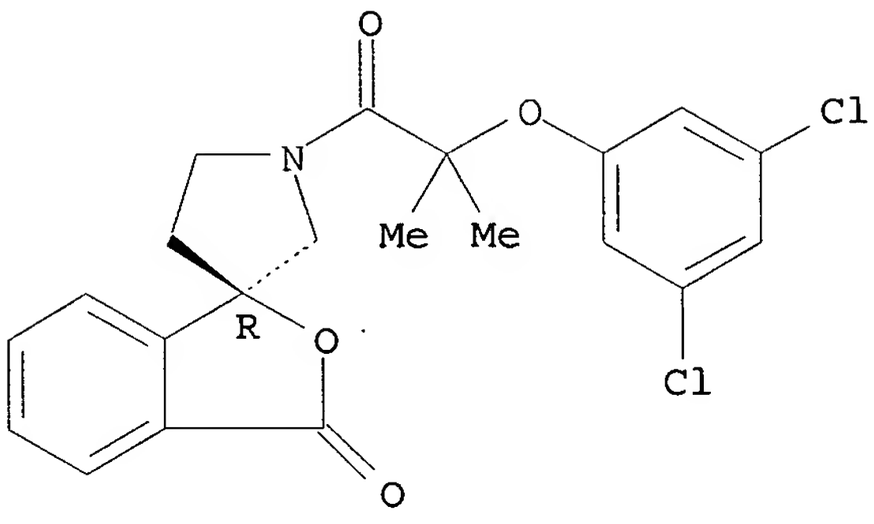
RN 872986-15-7 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-(2-methyl-1-oxo-2-phenoxypropyl)-, (1R)- (CA INDEX NAME)

Absolute stereochemistry.



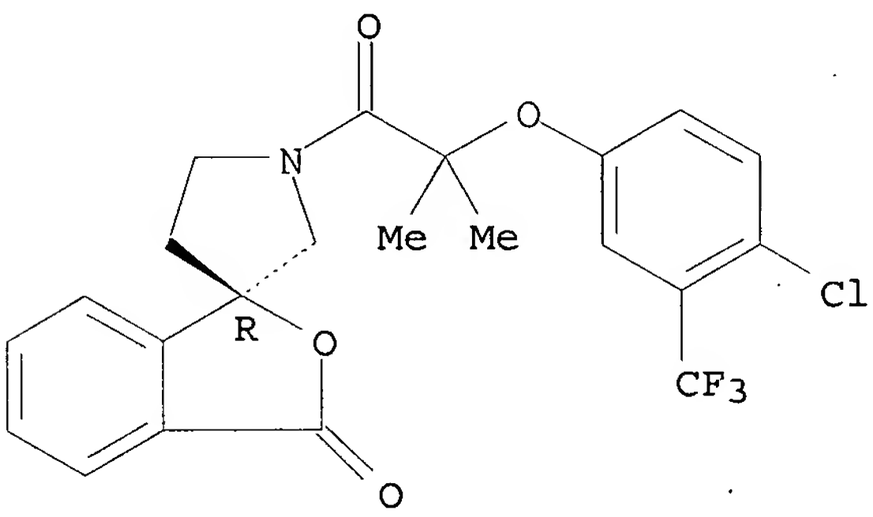
RN 872986-19-1 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(3,5-dichlorophenoxy)-2-methyl-1-oxopropyl]-, (1R)- (CA INDEX NAME)

Absolute stereochemistry.



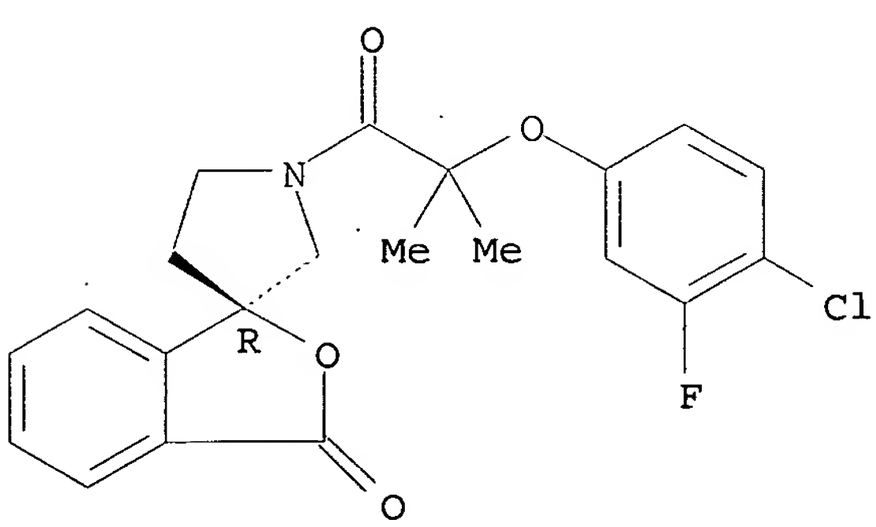
RN 872986-21-5 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-[4-chloro-3-(trifluoromethyl)phenoxy]-2-methyl-1-oxopropyl]-, (1R)- (CA INDEX NAME)

Absolute stereochemistry.



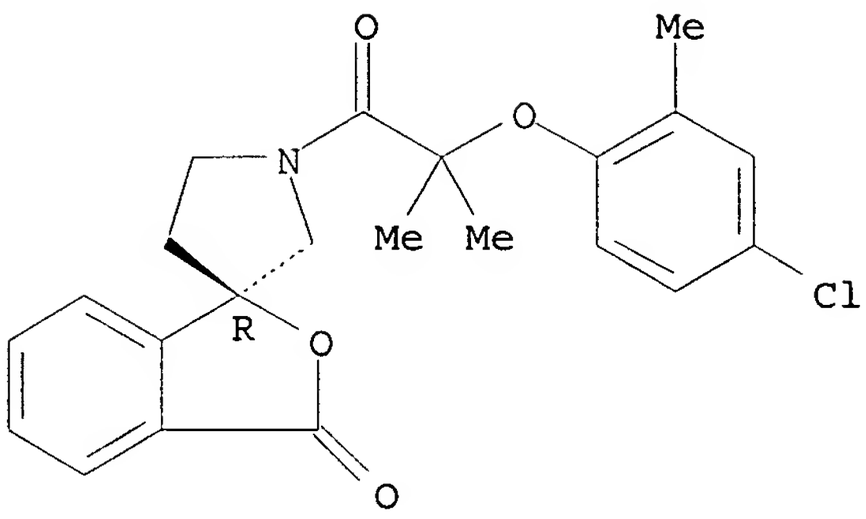
RN 872986-23-7 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(4-chloro-3-fluorophenoxy)-2-methyl-1-oxopropyl]-, (1R)- (CA INDEX NAME)

Absolute stereochemistry.



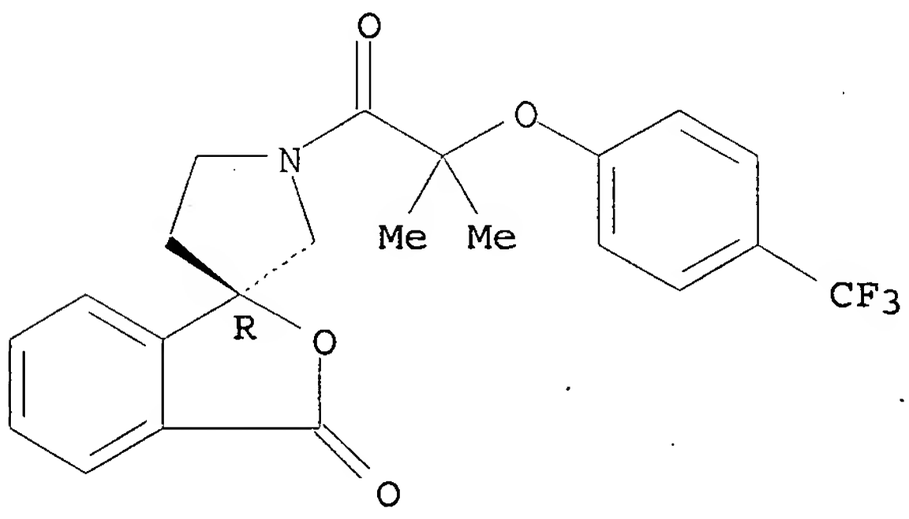
RN 872986-25-9 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-(4-chloro-2-methylphenoxy)-2-methyl-1-oxopropyl]-, (1R)- (CA INDEX NAME)

Absolute stereochemistry.



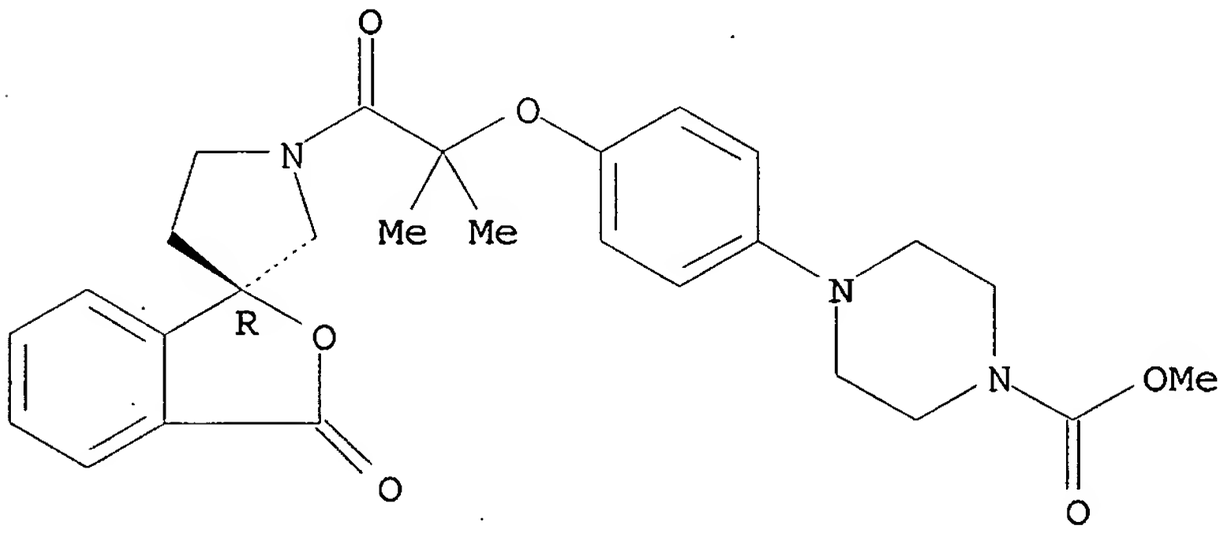
RN 872986-27-1 CAPLUS
CN Spiro[isobenzofuran-1(3H),3'-pyrrolidin]-3-one, 1'-[2-methyl-1-oxo-2-[4-(trifluoromethyl)phenoxy]propyl]-, (1R)- (CA INDEX NAME)

Absolute stereochemistry.



RN 872986-38-4 CAPLUS
CN 1-Piperazinecarboxylic acid, 4-[4-[1,1-dimethyl-2-oxo-2-[(1R)-3-oxospiro[isobenzofuran-1(3H),3'-pyrrolidin]-1'-yl]ethoxy]phenyl]-, methyl ester (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2005:588965 CAPLUS
DOCUMENT NUMBER: 143:115452
TITLE: Preparation of tricyclic spiropiperidines as

modulators of chemokine receptor activity
INVENTOR(S): Hossain, Nafizal; Ivanova, Svetlana
PATENT ASSIGNEE(S): Astrazeneca AB, Swed.
SOURCE: PCT Int. Appl., 70 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005061499	A1	20050707	WO 2004-SE1938	20041220
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RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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AU 2004303735	B2	20070920		
CA 2548494	A1	20050707	CA 2004-2548494	20041220
EP 1699791	A1	20060913	EP 2004-809111	20041220
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK, HR, IS				
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CN 1918160	A	20070221	CN 2004-80042013	20041220
JP 2007515476	T	20070614	JP 2006-546906	20041220
MX 2006PA07025	A	20060831	MX 2006-PA7025	20060619
US 2007099945	A1	20070503	US 2006-583468	20060620
IN 2006MN00848	A	20070518	IN 2006-MN848	20060718
NO 2006003355	A	20060922	NO 2006-3355	20060719
PRIORITY APPLN. INFO.:			SE 2003-3541 A	20031222
			WO 2004-SE1938 W	20041220

OTHER SOURCE(S): CASREACT 143:115452; MARPAT 143:115452
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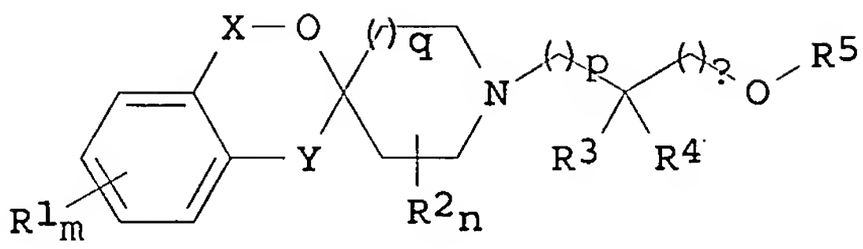

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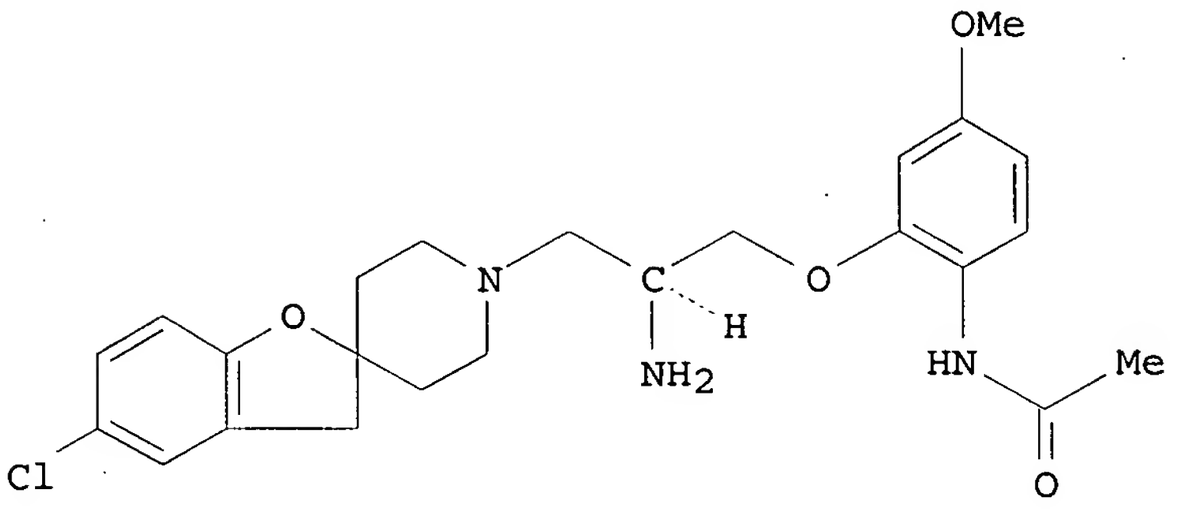


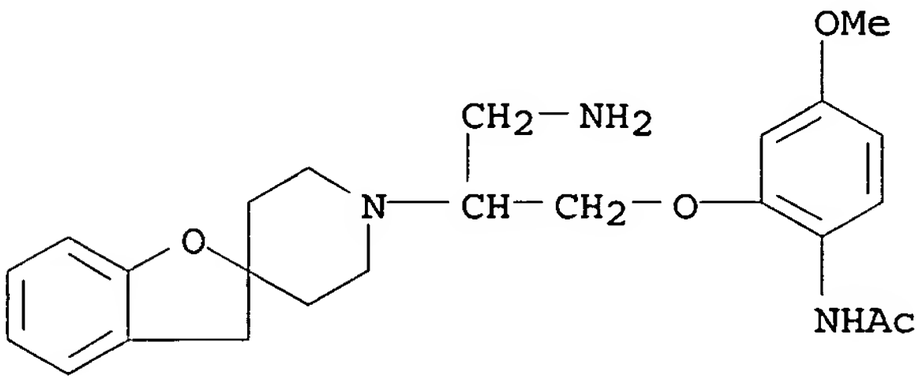
II

AB Title compds. I [m = 0-4; R1 = halo, CN, OH, etc.; X = bond, CH2 and Y = bond, CH2 provided that X, Y do not both simultaneously represent bond, CH2; n = 0-2; R2 = halo, alkyl, haloalkyl; q = 0-1; p = 0-2; R3 = halo, amino, carboxyl, etc.; R4 = H, alkyl, haloalkyl, halo; a = 0-2 provided that p and a are not both 0; R5 = (un)saturated 5-10-membered ring system] are prepared For instance, II is prepared in 4 steps from 5-methoxy-2-nitrophenol, (S)-oxiran-2-ylmethanol, and 5-chlorospiro[3H-benzofuran-2,4'-piperidine] (preparation given). I are modulators of chemokine receptor activity [no data] and useful for the treatment of, e.g., rheumatoid arthritis.

IT 857264-51-8P, N-[2-[3-Amino-2-(spiro[3H-benzofuran-2,4'-piperidin]-1'-yl)propoxy]-4-methoxyphenyl]acetamide
RL: BYP (Byproduct); PREP (Preparation)
(preparation of tricyclic spiropiperidines as modulators of chemokine receptor activity)

RN 857264-51-8 CAPLUS

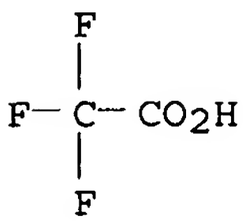
CN Acetamide, N-[2-(3-amino-2-spiro[benzofuran-2(3H),4'-piperidin]-1'-ylpropoxy)-4-methoxyphenyl]- (9CI) (CA INDEX NAME)



IT 857264-43-8P, N-[2-[[3-Amino-2-(5-chlorospiro[3H-benzofuran-2,4'-piperidin]-1'-yl)propyl]oxy]-4-methoxyphenyl]acetamide
RL: BYP (Byproduct); SPN (Synthetic preparation); PREP (Preparation)
(preparation of tricyclic spiropiperidines as modulators of chemokine receptor activity)

RN 857264-43-8 CAPLUS

CN Acetamide, N-[2-[3-amino-2-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)propoxy]-4-methoxyphenyl]- (CA INDEX NAME)



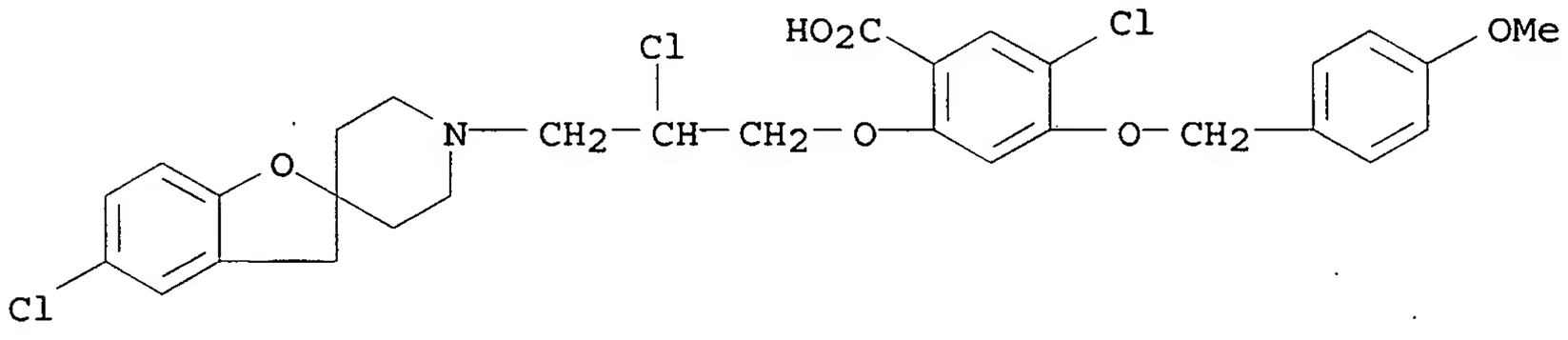
IT 857264-79-0P, 5-Chloro-2-[2-chloro-3-(5-chlorospiro[3H-benzofuran-2,4'-piperidin]-1'-yl)propoxy]-4-[(4-methoxybenzyl)oxy]benzoic acid trifluoroacetate
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of tricyclic spiropiperidines as modulators of chemokine receptor activity)

RN 857264-79-0 CAPLUS

CN Benzoic acid, 5-chloro-2-[2-chloro-3-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)propoxy]-4-[(4-methoxyphenyl)methoxy]-, trifluoroacetate (9CI) (CA INDEX NAME)

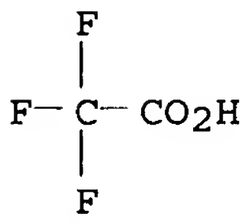
CM 1

CRN 857264-78-9
CMF C30 H30 Cl3 N O6



CM 2

CRN 76-05-1
CMF C2 H F3 O2



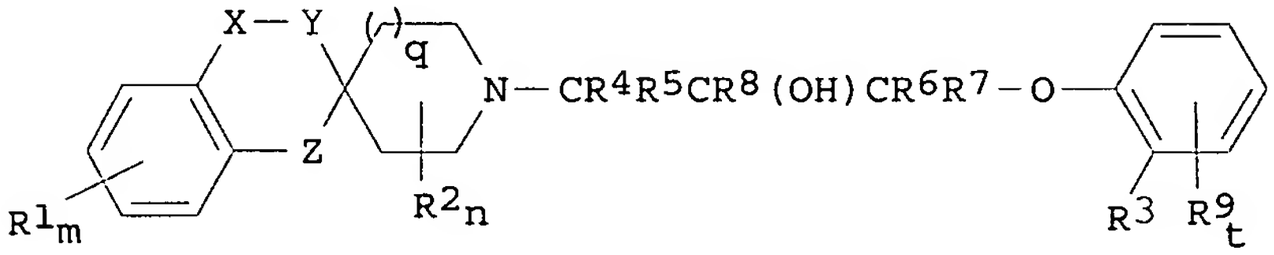
REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:41477 CAPLUS
DOCUMENT NUMBER: 140:93937
TITLE: Preparation of tricyclic spiropiperidines or spiropyrrolidines useful against disorders affected by modulation of chemokine receptors
INVENTOR(S): Hossain, Nafizal; Ivanova, Svetlana; Mensonides-Harsema, Marguerite
PATENT ASSIGNEE(S): Astrazeneca AB, Swed.
SOURCE: PCT Int. Appl., 281 pp.

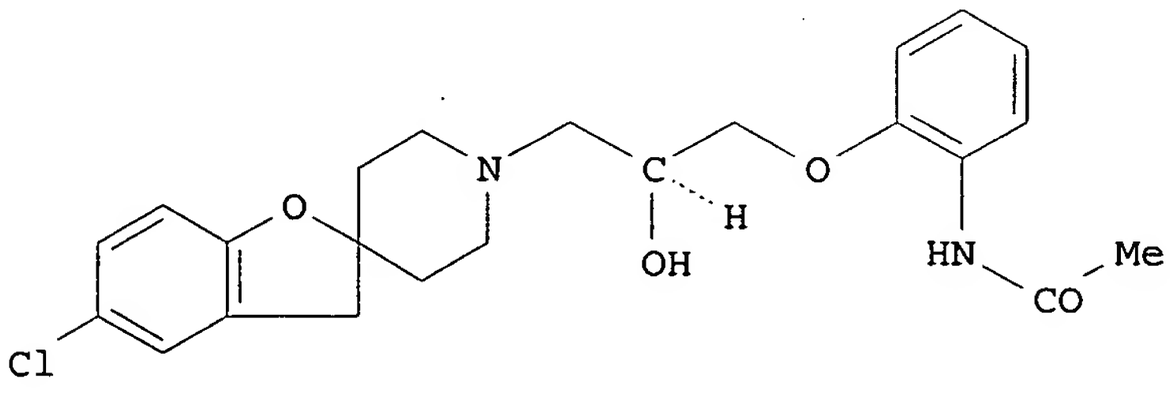
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
WO 2004005295	A1	20040115	WO 2003-SE1185		20030707
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW					
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG					
CA 2492122	A1	20040115	CA 2003-2492122		20030707
AU 2003243122	A1	20040123	AU 2003-243122		20030707
EP 1521757	A1	20050413	EP 2003-762957		20030707
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK					
BR 2003012560	A	20050510	BR 2003-12560		20030707
CN 1675218	A	20050928	CN 2003-819146		20030707
JP 2005537255	T	20051208	JP 2004-519472		20030707
NZ 537259	A	20060831	NZ 2003-537259		20030707
CN 1974574	A	20070606	CN 2006-10143556		20030707
IN 2004DN04014	A	20070427	IN 2004-DN4014		20041216
ZA 2005000024	A	20060222	ZA 2005-24		20050103
MX 2005PA00278	A	20050331	MX 2005-PA278		20050104
US 2005245741	A1	20051103	US 2005-520699		20050107
NO 2005000635	A	20050331	NO 2005-635		20050204
PRIORITY APPLN. INFO.:			SE 2002-2133	A	20020708
			CN 2003-819146	A3	20030707
			WO 2003-SE1185	W	20030707

OTHER SOURCE(S): MARPAT 140:93937
GI



I



II

AB The invention provides tricyclic spiropiperidines or spiropyrrolidines (shown as I; variables defined below; e.g. II), processes for their preparation, pharmaceutical compns. containing them and their use in therapy for

disorders affected by modulation of chemokine receptors (no data). For I: m is 0-4; each R1 = halogen, cyano, hydroxy, C1-C6 alkyl, C1-C6 haloalkyl, C1-C6 alkoxy or sulfonamido; either X = a bond, -CH2-, -O- or -C(O)- and Y = a bond, -CH2-, -O- or -C(O)-, or X and Y together = -CH:CMe- or -CMe:CH-, and Z = a bond, -O-, -NH- or -CH2-, provided that only one of X, Y and Z can be a bond at any one time and provided that X and Y do not both simultaneously = -O- or -C(O)-. N = 0-2; each R2 = halogen or C1-C6 alkyl; q = 0-1; R3 = -NHC(O)R10, -C(O)NR11R12 or -COOR12a; R4, R5, R6, R7 and R8 = H or a C1-C6 alkyl group; t = 0-2; each R9 = halogen, cyano, hydroxy, carboxy, C1-C6 alkoxy, C1-C6 alkoxycarbonyl, C1-C6 haloalkyl, or C1-C6 alkyl; addnl. details are given in the claims. Methods of preparation are claimed and >200 example prepns. are included. For example, II was prepared in 2 steps starting from N-(2-hydroxyphenyl)acetamide, ((2S)-oxiran-2-yl)methyl and Cs2CO3 in DMF to give N-[2-[((2S)-oxiran-2-yl)methoxy]phenyl]acetamide as an intermediate, which was reacted with 5-chloro-3H-spiro[1-benzofuran-2,4'-piperidine] in EtOH to give II.

IT 644968-87-6P 644969-01-7P 644969-11-9P
644969-20-0P 644969-46-0P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of tricyclic spiropiperidines or spiropyrrolidines useful against disorders affected by modulation of chemokine receptors)

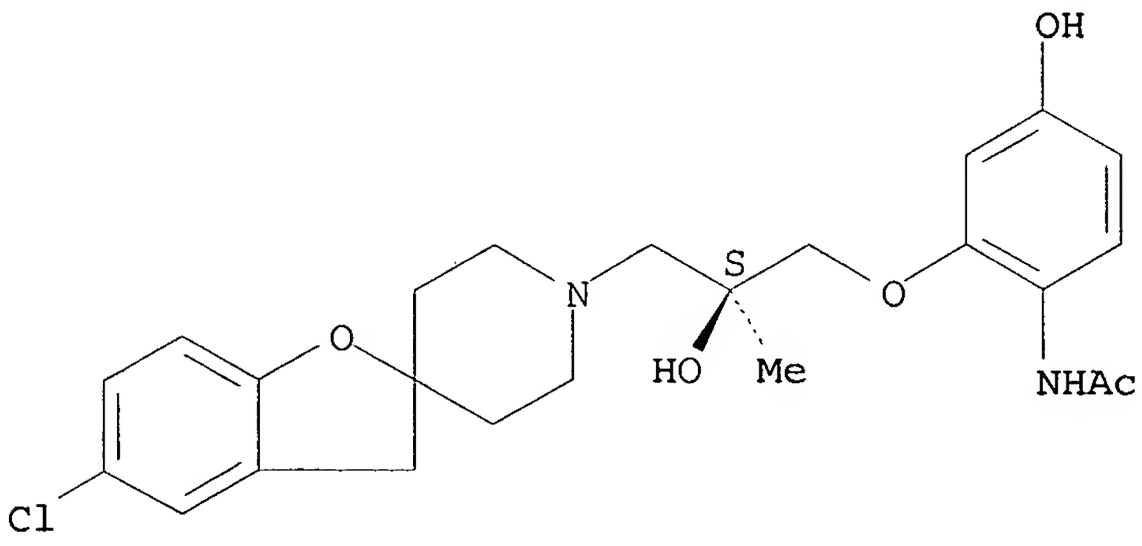
RN 644968-87-6 CAPLUS

CN Acetamide, N-[2-[(2S)-3-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-4-hydroxyphenyl]-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

CM 1

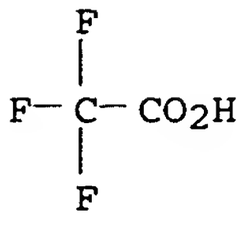
CRN 644968-86-5
CMF C24 H29 Cl N2 O5

Absolute stereochemistry.



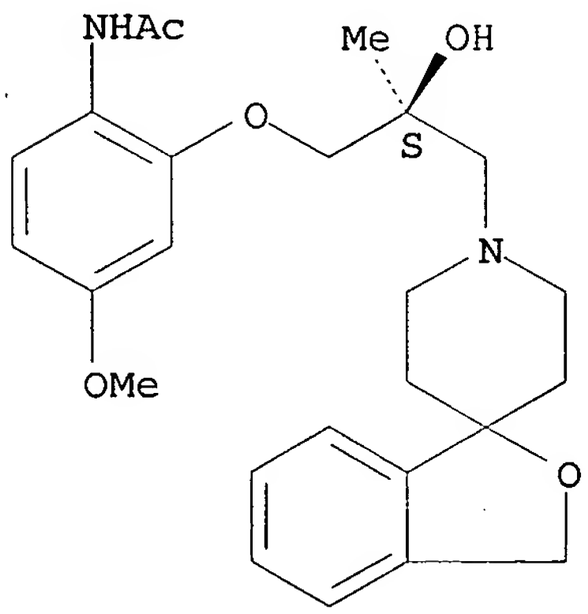
CM 2

CRN 76-05-1
CMF C2 H F3 O2



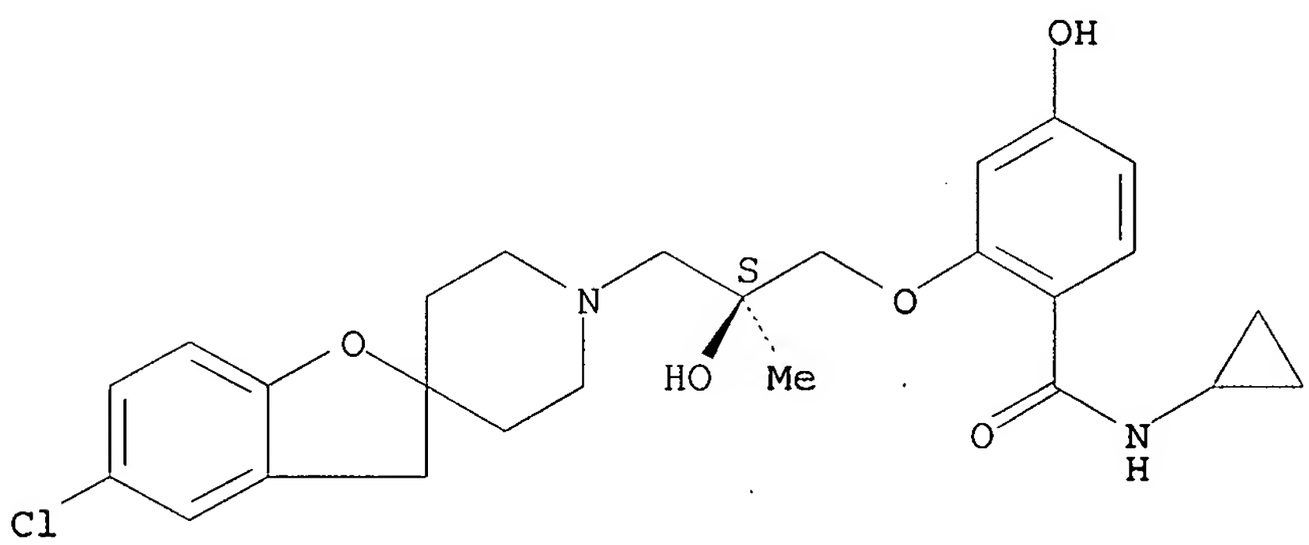
RN 644969-01-7 CAPLUS
CN Acetamide, N-[2-[(2S)-2-hydroxy-2-methyl-3-spiro[isobenzofuran-1(3H),4'-piperidin]-1'-ylpropoxy]-4-methoxyphenyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 644969-11-9 CAPLUS
CN Benzamide, 2-[(2S)-3-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-N-cyclopropyl-4-hydroxy- (CA INDEX NAME)

Absolute stereochemistry.

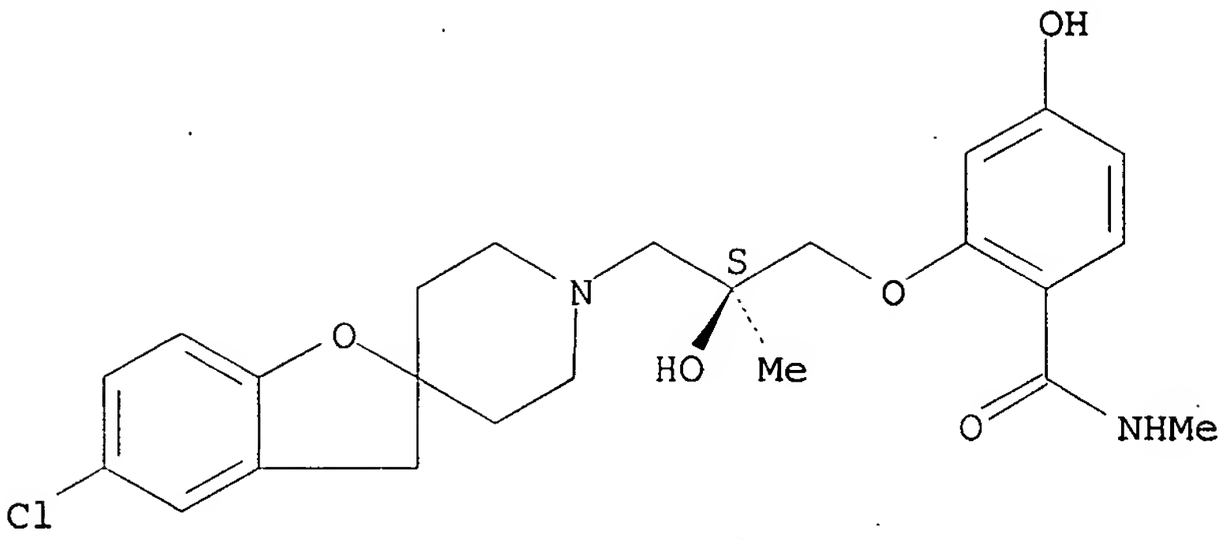


RN 644969-20-0 CAPLUS
CN Benzamide, 2-[(2S)-3-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-4-hydroxy-N-methyl-, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

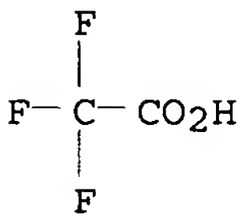
CM 1

CRN 644969-19-7
CMF C24 H29 Cl N2 O5

Absolute stereochemistry.

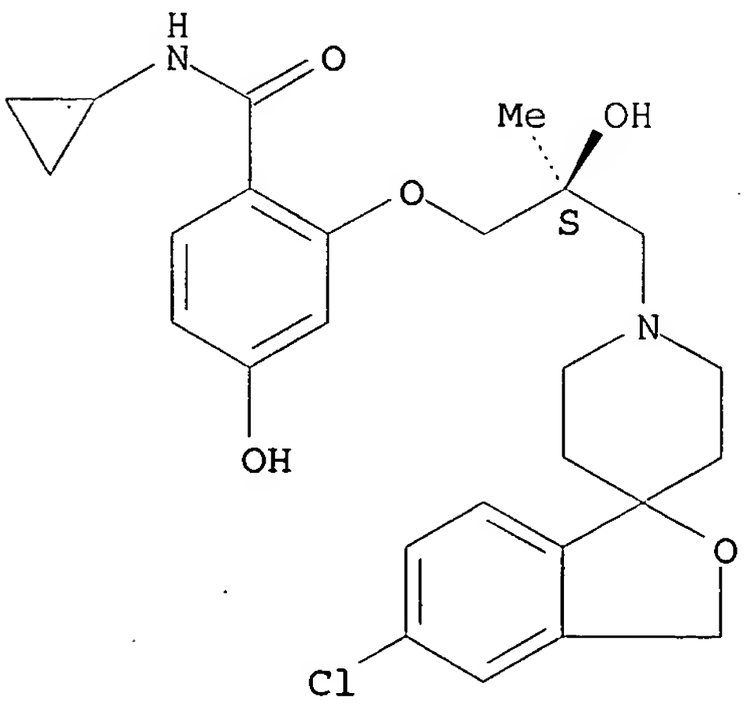


CM 2
CRN 76-05-1
CMF C2 H F3 O2



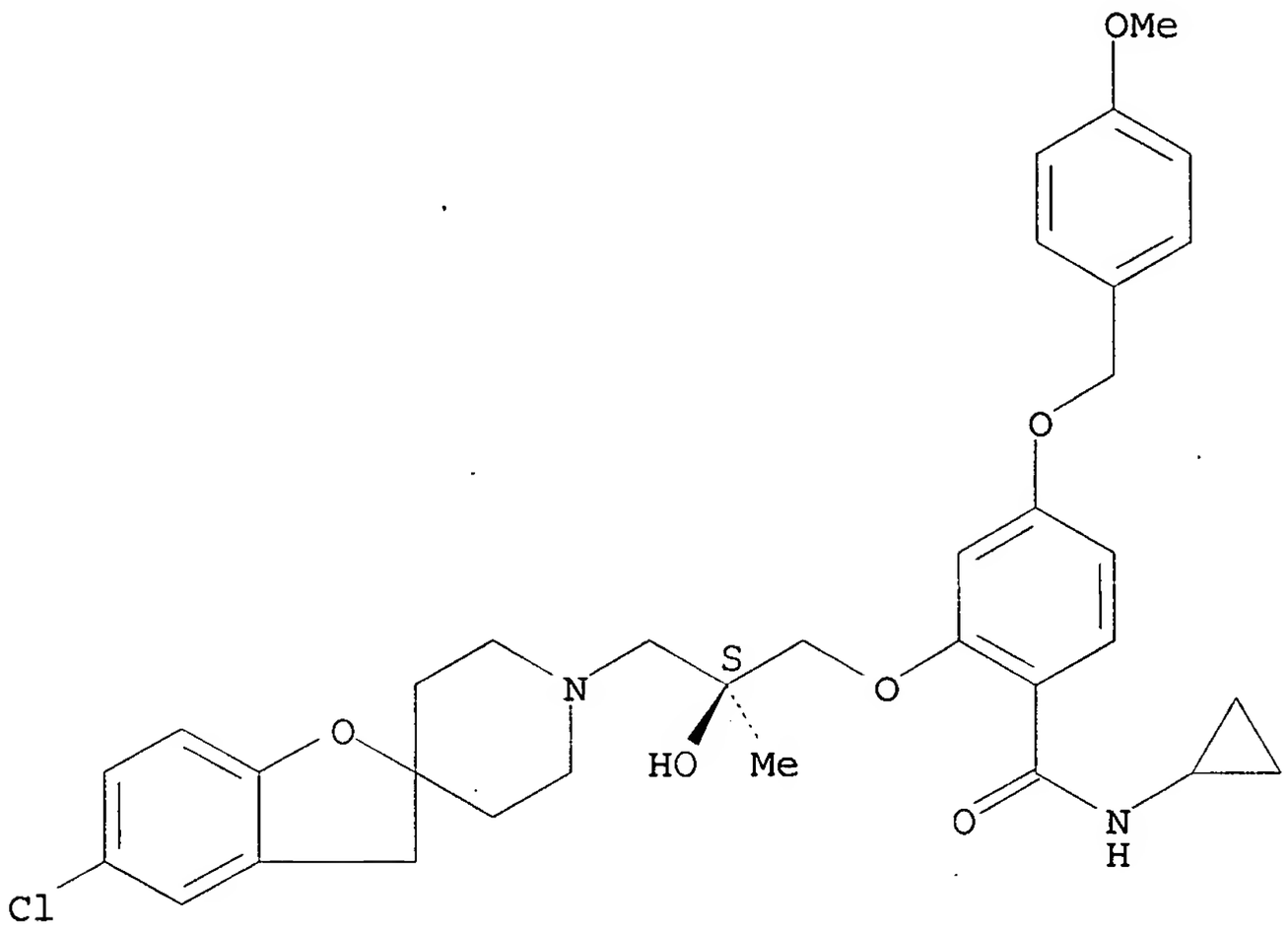
RN 644969-46-0 CAPLUS
CN Benzamide, 2-[(2S)-3-(5-chlorospiro[isobenzofuran-1(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-N-cyclopropyl-4-hydroxy- (CA INDEX NAME)

Absolute stereochemistry.



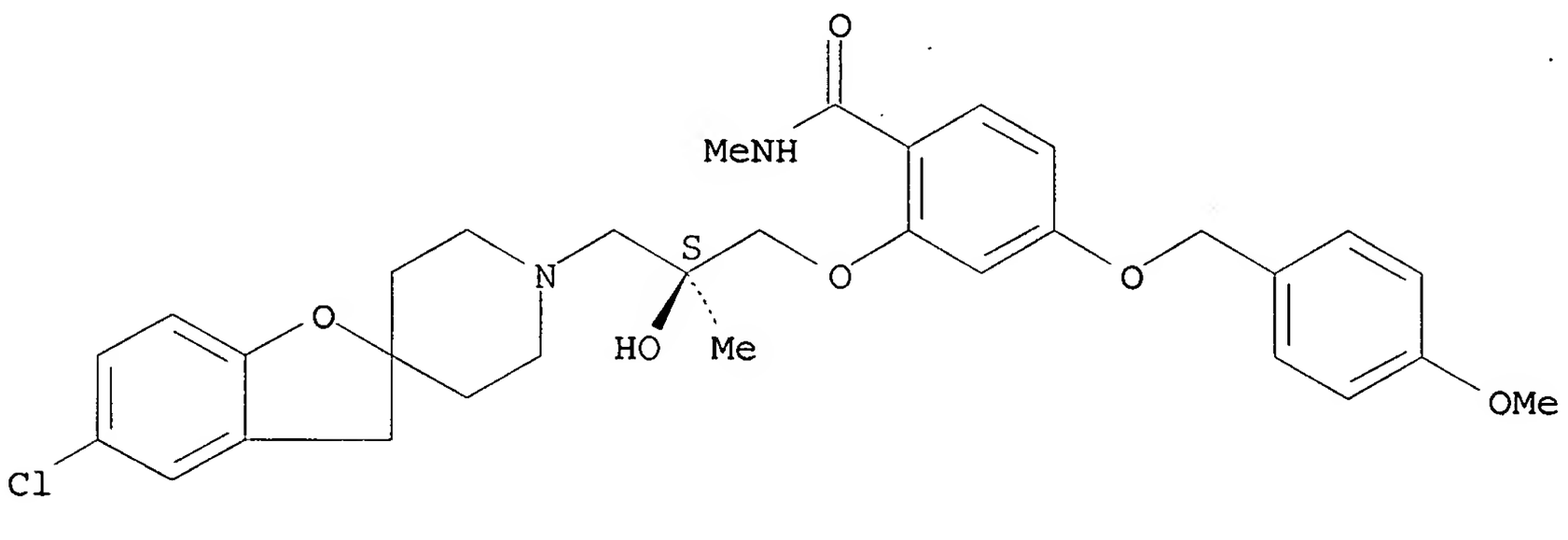
IT 644969-14-2P 644969-23-3P 644969-47-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of tricyclic spiropiperidines or spiropyrrolidines useful against disorders affected by modulation of chemokine receptors)
RN 644969-14-2 CAPLUS
CN Benzamide, 2-[(2S)-3-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-N-cyclopropyl-4-[(4-methoxyphenyl)methoxy]- (CA INDEX NAME)

Absolute stereochemistry.



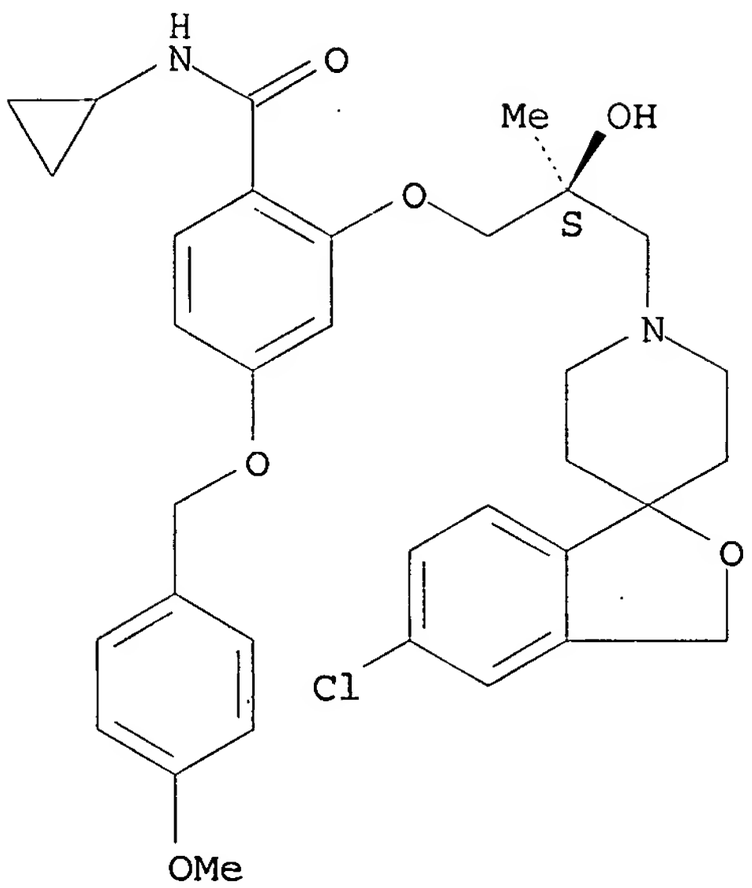
RN 644969-23-3 CAPLUS
CN Benzamide, 2-[(2S)-3-(5-chlorospiro[benzofuran-2(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-4-[(4-methoxyphenyl)methoxy]-N-methyl- (CA INDEX NAME)

Absolute stereochemistry.



RN 644969-47-1 CAPLUS
CN Benzamide, 2-[(2S)-3-(5-chlorospiro[isobenzofuran-1(3H),4'-piperidin]-1'-yl)-2-hydroxy-2-methylpropoxy]-N-cyclopropyl-4-[(4-methoxyphenyl)methoxy]- (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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FILE 'REGISTRY' ENTERED AT 11:44:10 ON 11 JAN 2008
L1 STRUCTURE UPLOADED
L2 2 S L1
L3 65 S L1 FULL

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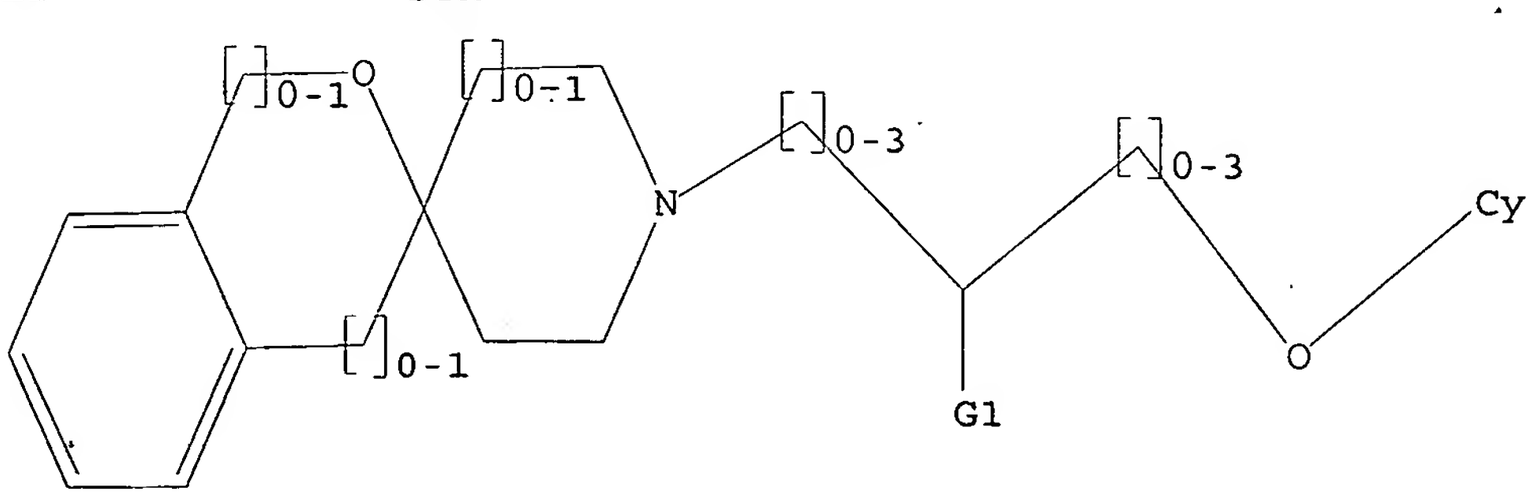
L4 3 S L3

=> d l1
L1 HAS NO ANSWERS
L1 STR



G1 C,N,X

Structure attributes must be viewed using STN Express query preparation.

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